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Tanabe et al.

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[54] RESISTOR PASTE AND CERAMIC SUBSTRATE

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[21] Appl. No.: **899,156**

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[57] ABSTRACT

Related U.S. Application Data

[63] Continuation of Ser. No. 531,750, Jun. 1, 1990, abandoned.

A resistor paste comprising an inorganic component which consists essentially of from 20 to 70 wt % of glass powder and from 30 to 80 wt % of a powder selected from the group consisting of SnO₂ powder, Sb-doped SnO₂ powder and a mixture thereof, wherein the glass powder consists essentially of from 12 to 50 wt % of SiO₂, from 0 to 20 wt % of Al₂O₃, from 0 to 40 wt % of MgO, from 0 to 40 wt % of CaO, from 0 to 60 wt % of SrO, from 16 to 60 wt % of MgO+CaO+SrO, from 0 to 10 wt % of Li₂O+Na₂O+K₂O+Cs₂O, from 0 to 10 wt % of PbO, from 0 to 20 wt % of ZnO, from 0 to 10 wt % of ZrO₂+TiO₂, from 8 to 40 wt % of B₂O₃, from 0 to 60 wt % of Ta₂O₅, from 0 to 50 wt % of Nb₂O₅, from 0 to 60 wt % of Ta₂O₅+Nb₂O₅, and from 0.1 to 20 wt % of Fe₂O₃+CuO+NiO+CoO+MnO+MoO₃+WO₃+Cr₂O₃+Bi₂O₃+CeO₂+Sb₂O₃+In₂O₃+SnO₂.

[30] Foreign Application Priority Data

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Jul. 14, 1989 [JP] Japan 1-180565

[51] Int. Cl.⁵ **C03C 8/14**

[52] U.S. Cl. **501/17; 501/32; 501/66; 501/78**

[58] Field of Search 501/17, 32, 64, 65, 501/72, 73, 77, 78

[56] References Cited

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3 Claims, 3 Drawing Sheets

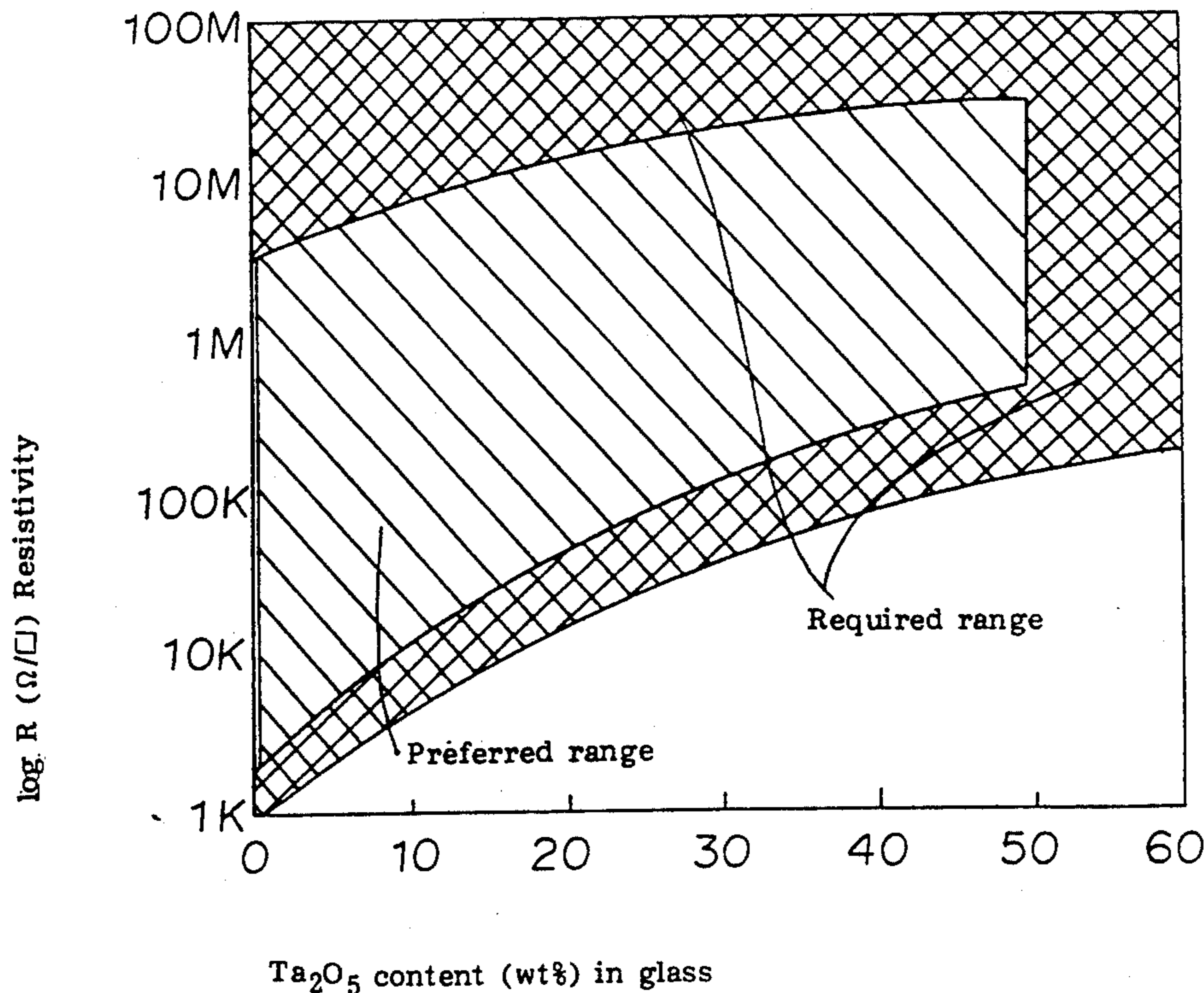


FIGURE 1

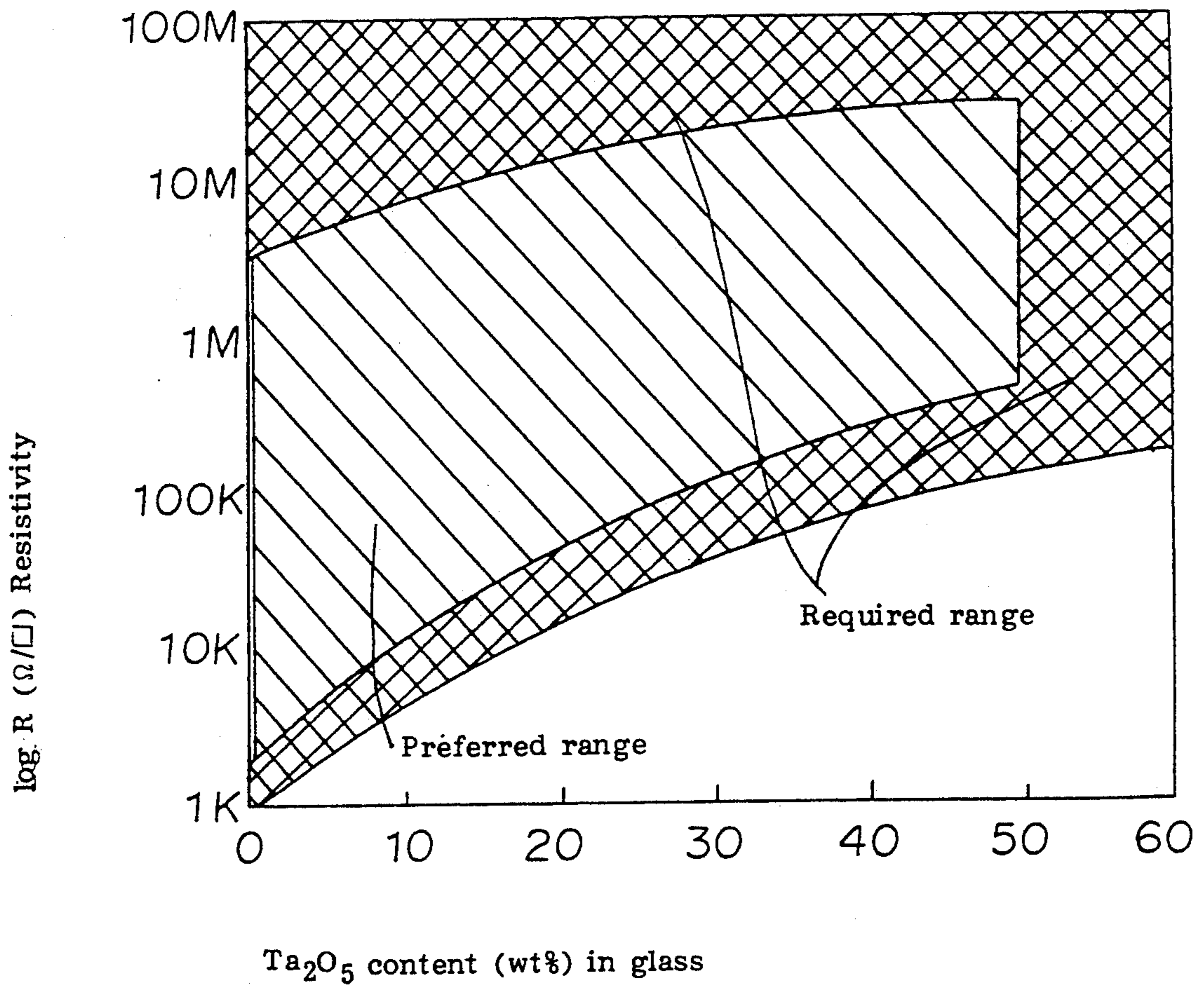


FIGURE 2

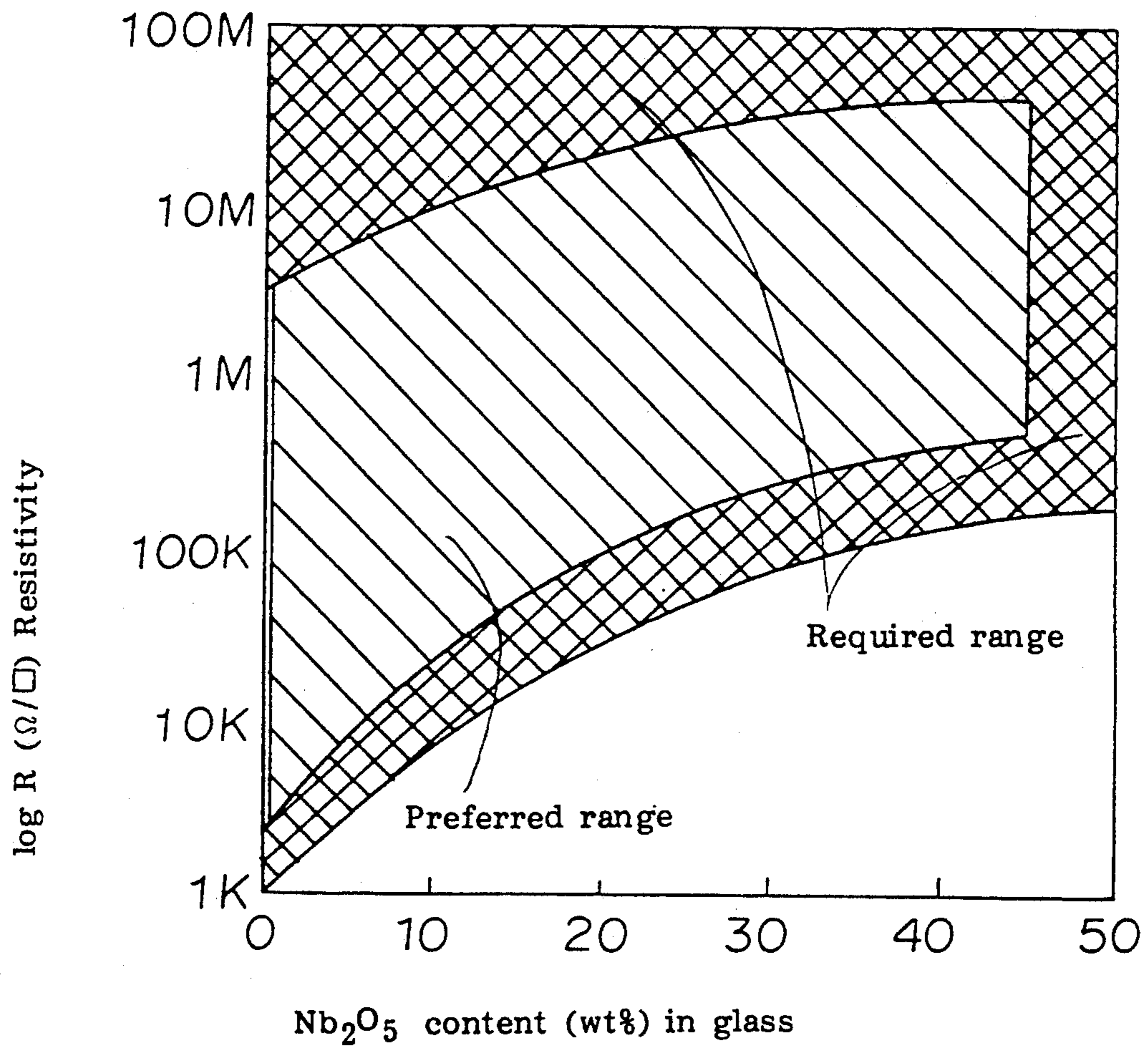
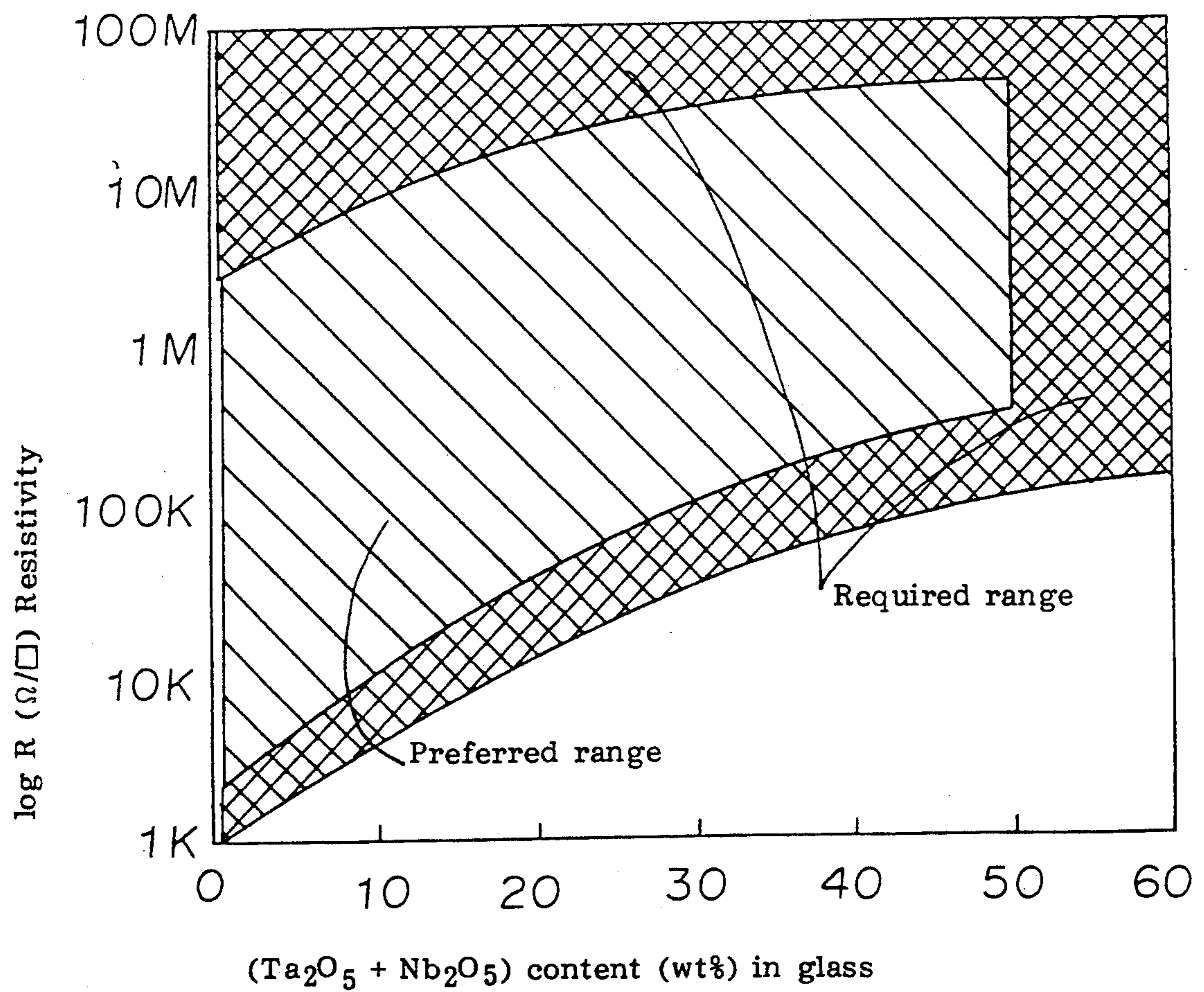


FIGURE 3



RESISTOR PASTE AND CERAMIC SUBSTRATE

This application is a continuation of application Ser. No. 07/531,750, filed on Jun. 1, 1990, now abandoned. 5

BACKGROUND OF THE INVENTION

1. Field of the Invention

The present invention relates to a resistor paste composition and a ceramic substrate. More particularly, it relates to a resistor paste comprising $\text{SiO}_2\text{-B}_2\text{O}_3$ system glass containing metal oxides, and SnO_2 and/or Sb-doped SnO_2 , which is useful for a ceramic substrate, whereby even when fired in a non-oxidizing atmosphere, it will not be reduced, and provides adequate stability of e.g. resistivity. 15

2. Discussion of Background

Heretofore, the resistor in a hybrid integrated circuit has been formed in such a manner that a silver (Ag) or Ag-palladium (Pd) conductor is formed on or in a ceramic substrate, and a resistor paste is printed between the conductor patterns, followed by firing in an oxidizing atmosphere such as air at a temperature of from about 850° to 900° C. The resistor paste used in such a case used to be composed mainly of RuO_2 and glass. However, recently, it has been common to employ a copper (Cu) conductor instead of the Ag or Ag-Pd conductor, from the viewpoint of the reliability in e.g. migration. 25

However, the Cu conductor is oxidized unless it is fired in a non-oxidizing atmosphere such as nitrogen. In a non-oxidizing atmosphere, RuO_2 can not be used, since it will be reduced and will not form a resistor in such a non-oxidizing atmosphere. 30

Therefore, it has recently been proposed to employ a combination of antimony (Sb)-doped tin oxide and glass powder (Japanese Unexamined Patent Publication No. 119902/1987). However, such a combination still has a drawback that the resistivity and the temperature coefficient of resistivity (TCR) are not yet adequately stable. 40

SUMMARY OF THE INVENTION

It is an object of the present invention to provide a novel resistor paste and ceramic substrate, which can be fired in a non-oxidizing atmosphere such as nitrogen and which are able to provide a constant resistivity and temperature coefficient of resistivity (TCR). 45

The present invention has been made to solve the above-mentioned problems and provides a resistor paste comprising an inorganic component which consists essentially of from 20 to 70 wt % of glass powder and from 30 to 80 wt % of a powder selected from the group consisting of SnO_2 powder, Sb-doped SnO_2 powder and a mixture thereof, wherein the glass powder consists essentially of from 12 to 50 wt % of SiO_2 , from 0 to 20 wt % of Al_2O_3 , from 0 to 40 wt % of MgO , from 0 to 40 wt % of CaO , from 0 to 60 wt % of SrO , from 16 to 60 wt % of MgO CaO+SrO , from 0 to 10 wt % of $\text{Li}_2\text{O+Na}_2\text{O+K}_2\text{O+Cs}_2\text{O}$, from 0 to 10 wt % of PbO , from 0 to 20 wt % of ZnO , from 0 to 10 wt % of $\text{ZrO}_2+\text{TiO}_2$, from 8 to 40 wt % of B_2O_3 , from 0 to 60 wt % of Ta_2O_5 , from 0 to 50 wt % of Nb_2O_5 , from 0 to 60 wt % of $\text{Ta}_2\text{O}_5+\text{Nb}_2\text{O}_5$, and from 0.1 to 20 wt % of $\text{Fe}_2\text{O}_3+\text{CuO+NiO+CoO+MnO+MoO}_3+\text{WO}_3+\text{Cr}_2\text{O}_3+\text{Bi}_2\text{O}_3+\text{CeO}_2+\text{Sb}_2\text{O}_3+\text{In}_2\text{O}_3+\text{SnO}_2$. 55

The present invention also provides a ceramic substrate having such a resistor paste fired thereon. 65

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a graph showing the required range and the preferred range of the amount of Ta_2O_5 relative to the resistivity in the case where Ta_2O_5 is used alone ($\text{Nb}_2\text{O}_5 < 0.1$ wt %).

FIG. 2 is a graph showing the required range and the preferred range of the amount of Nb_2O_5 relative to the resistivity in the case where Nb_2O_5 is used alone ($\text{Ta}_2\text{O}_5 < 0.1$ wt %).

FIG. 3 is a graph showing the required range and the preferred range of the amount of $\text{Ta}_2\text{O}_5+\text{Nb}_2\text{O}_5$ relative to the resistivity.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Now, the present invention will be described in detail.

The resistor paste of the present invention is suitable for use for a single layer or multi-layer ceramic substrate, and is useful in such a manner that it is formed by a method of e.g. printing, on a ceramic substrate such as a fired and solidified alumina substrate, or on a green sheet for a ceramic substrate, followed by firing in a non-oxidizing atmosphere such as a nitrogen atmosphere. In this specification, “%” means “wt %” unless otherwise specified. In the resistor paste of the present invention, the inorganic component consists essentially of from 20 to 70% of glass powder and from 30 to 80% of a powder of conductive material. These materials will be described below.

The glass powder is preferably a sinterable $\text{SiO}_2\text{-B}_2\text{O}_3$ system glass which has adequate fluidity at a low temperature (e.g. at a temperature of not higher than 900° C.) and which is capable of covering and sufficiently wetting the above powder of conductive material at the time of firing.

If the content of such glass powder is less than 20%, the powder of conductive material can not adequately be wetted, whereby the sintered layer will have void pores, and strength of the resistor formed by firing the resistor paste of the present invention will be low, and the stability of the resistivity will be low. On the other hand, if it exceeds 70%, the adhesion among the powder particles of conductive material tends to be small, whereby the resistivity tends to be high.

The glass powder in the present invention is preferably within a range of from 25 to 65% within the above range.

On the other hand, as the powder of conductive material, commercially available SnO_2 or SnO_2 having Sb doped usually in the form of an oxide of Sb_2O_3 , may be used alone or in combination, because such material has high conductivity i.e. low resistivity, so that the resistivity of the resistor of the present invention which is a composite of the conductive material and the glass, can be adjusted to a desired level.

Sb-doped SnO_2 has a low resistivity as compared with non-doped SnO_2 . The resistivity increases if the doping amount increases to excess. When the resistance according to the present invention is not higher than 10 $\text{M}\Omega$, the doping amount is suitably within a range of from 0 to 20%, preferably from 0.1 to 15%, more preferably from 1 to 10%, as calculated as an oxide of Sb_2O_3 . When the resistance according to the present invention is higher than 10 $\text{M}\Omega$, the doping amount may be 20% or higher as calculated as an oxide of Sb_2O_3 .

With respect to the particle size of the glass powder according to the present invention, if the particle size is too small, the above resistivity tends to be too high, and if it is too large, it becomes difficult to adequately wet the glass, and void pores tend to increase in the sintered layer, such being undesirable. The average particle size should usually be within a range of from 0.5 to 6 μm , preferably from 1 to 5 μm .

On the other hand, if the particle size of the powder of conductive material according to the present invention, is too small, the resistivity tends to be too large, and if it is too large, the distribution on the ceramic substrate tends to be non-uniform, and the variation in the resistivity will be large, such being undesirable. The average particle size should usually be within a range of from 0.01 to 5 μm , preferably from 0.05 to 3 μm .

In the present invention, the glass powder consists essentially of:

SiO ₂	12-50%
Al ₂ O ₃	0-20%
MgO + CaO + SrO (MgO, 0-40%, CaO 0-40%, SrO 0-60%)	16-60%
Li ₂ O + Na ₂ O + K ₂ O + Cs ₂ O	0-10%
PbO	0-10%
ZnO	0-20%
ZrO ₂ + TiO ₂	0-10%
B ₂ O ₃	8-40%
Ta ₂ O ₅	0-60%
Nb ₂ O ₅	0-50%
Ta ₂ O ₅ + Nb ₂ O ₅	0-60%
Metal oxides	0.1-20%

These components will be described below.

In such a composition, SiO₂ is a network former of the glass, and if it is less than 12%, the softening point will be too low, whereby the heat resistance will be low, and the glass will easily be deformed when it is fired again, such being undesirable. On the other hand, if SiO₂ exceeds 50%, the softening point tends to be too high, whereby the fluidity of glass tends to be poor at the time of firing, and it tends to be incapable of covering and wetting the powder of conductive material. Further, void pores in the sintered layer tend to be too many, and the stability of the resistance will be poor. It is preferably within a range of from 15 to 45%.

Al₂O₃ is not essential, but when incorporated, it contributes to the improvement of moisture resistance. If it exceeds 20%, the softening temperature of glass will be high, and the sinterability tends to be poor. It is preferably not higher than 18%.

MgO+CaO+SrO improve the solubility at the time of the preparation of the glass powder and thus have a function to adjust the thermal expansion coefficient. If their content is less than 16%, the above solubility will not adequately be improved, and devitrification is likely to result during the preparation of glass. On the other hand, if it exceeds 60%, the thermal expansion coefficient will be large, such being undesirable. Preferably, it is within a range of from 18 to 55%.

Further, in the above MgO+CaO+SrO, if either MgO or CaO is 40% or higher, the thermal expansion coefficient tends to be too large. Preferably, it is within a range of from 0 to 35%. In the above MgO+CaO+SrO, if SrO is 60% or higher, the thermal expansion coefficient tends to be too large. Preferably, it is within a range of from 0 to 55%.

Li₂O+Na₂O+K₂O+Cs₂O are not essential, but they are effective for the improvement of the solubility of glass and also have a function to increase the resistivity.

If their content exceeds 10%, the thermal expansion coefficient tends to be too large, the matching with the substrate will be poor, and the possibility of cracking in a thick film after firing increases. Preferably, it is not more than 8%.

PbO is not essential, but it is effective as a flux component for glass and has a function to increase the resistivity. If it exceeds 10%, the resistivity tends to be unstable. Preferably, it is not more than 5%.

ZnO is not essential, but may be incorporated up to 20% in order to improve the solubility of glass. Preferably, it is not more than 15%.

ZrO₂+TiO₂ are not essential. However, when they are incorporated, the moisture resistance of the resistor may be improved. It may be added up to 10%, preferably not more than 7%.

B₂O₃ is used as a flux component. If it is less than 8%, the softening point will be high, the sintering tends to be inadequate, and void pores in the sintered layer tend to be too many. On the other hand, if it exceeds 40%, the water resistance of glass will be low. Preferably, it is within a range of from 10 to 38%.

Ta₂O₅ and Nb₂O₅ are not essential components, but they are useful for adjusting the resistivity and the temperature coefficient of resistivity (TCR). By the incorporation of Ta₂O₅ and Nb₂O₅, it is possible to shift the resistivity to a higher direction and to shift TCR to the positive direction. Their amounts are determined to meet the desired resistivity. However, if Ta₂O₅ exceeds 60%, or Nb₂O₅ exceeds 50%, when manufacturing glass vitrification tends to be difficult.

The required ranges and the preferred ranges of Ta₂O₅, Nb₂O₅ and Ta₂O₅+Nb₂O₅ are shown in FIGS. 1, 2 and 3, respectively. The main points of these FIGS. 1 to 3 will be summarized as follows.

FIG. 1 is a graph showing the required range and the preferred range of the amount of Ta₂O₅ relative to the resistivity in a case where Ta₂O₅ is used alone (Nb₂O₅<0.1 wt %).

FIG. 2 is a graph showing the required range and the preferred range of the amount of Nb₂O₅ relative to the resistivity in a case where Nb₂O₅ is used alone (Ta₂O₅<0.1 wt %).

FIG. 3 is a graph showing the required range and the preferred range of the amount of Ta₂O₅+Nb₂O₅ relative to the resistivity.

Here, the combined use means that each of Ta₂O₅ and Nb₂O₅ is at least 0.1%. Namely, if one of them is less than 0.1%, the other is regarded as a single use.

Respective ranges of Ta₂O₅ and Nb₂O₅ for desired levels of resistivity (Either Ta₂O₅ or Nb₂O₅ is used alone)

Resistivity (Ω/\square)	Ta ₂ O ₅ (wt %)		Nb ₂ O ₅ (wt %)	
	Required range	Preferred range	Required range	Preferred range
10K	0-18	0.5-9	0-12	0.5-6
100K	0-46	0.5-30	0-34	0.5-20
1M	0-60	0.5-50	0-50	0.5-45
10M	0-60	15-50	0-50	10-45
100M	0-60	—	0-50	—

Respective ranges of Ta₂O₅ and Nb₂O₅ for desired levels of resistivity (Ta₂O₅ and Nb₂O₅ is used in combination)

Resistivity (Ω/\square)	Ta ₂ O ₅ + Nb ₂ O ₅ (wt %)	
	Required range	Preferred range
10K	0.2-18	0.5-9
100K	0.2-46	0.5-30
1M	0.2-60	0.5-50

-continued

10M	0.2-60	10-50
100M	0.2-60	—

As the above metal oxides, Fe₂O₃, CuO, NiO, MnO, MoO₃, WO₃, Bi₂O₃, CeO₂, CoO, Cr₂O₃, Sb₂O₃, In₂O₃ and SnO₂, may be used alone or in combination. These metal oxides have a function to adjust the resistivity and the temperature coefficient of resistivity (TCR) and to improve the laser trimming properties. Preferred among these metal oxides are NiO, MnO and Sb₂O₃. Particularly preferred is NiO. The respective functions will be listed below.

Fe₂O₃, CuO, NiO, MnO, CoO, Cr₂O₃, SnO₂, Sb₂O₃ and WO₃ are effective to lower the resistivity and shift TCR to the positive direction.

MoO₃ is effective to lower the resistivity and shift TCR to the negative direction.

CeO₂ is effective to increase the resistivity and shift TCR to the positive direction.

Bi₂O₃ is effective to increase the resistivity and shift TCR to the negative direction.

In₂O₃ is effective to increase the resistivity and shift TCR to the negative direction.

Further, they are effective to improve the cutting properties for the laser trimming to adjust the resistivity.

The respective amounts in the glass composition are determined to meet the desired resistance, temperature coefficient of resistivity (TCR) and laser trimming properties. However, if the total amount of the above metal oxides is less than 0.1%, no substantial effects will be obtained. On the other hand, if it exceeds 20%, the resistivity drift in the high temperature storage test will be large, such being undesirable. Preferably, the total amount is within a range of from 1 to 15%.

Among the above metal oxides, NiO, MnO and Sb₂O₃ are excellent in the effects for adjusting the resistivity and TCR and for stabilizing the resistivity drift. Among them, NiO is most excellent.

The preferred ranges with their resistivity being not more than 1M are as follows:

SiO ₂	15-45%
Al ₂ O ₃	0-18%
MgO + CaO + SrO (MgO, 0-35%, CaO 0-35%, SrO 0-35%)	18-55%
Li ₂ O + Na ₂ O + K ₂ O + Cs ₂ O	0-8%
PbO	0-5%
ZnO	0-15%
ZrO ₂ + TiO ₂	0-7%
B ₂ O ₃	10-38%
Ta ₂ O ₅	0.5-50%
Nb ₂ O ₅	0.5-45%
Ta ₂ O ₅ + Nb ₂ O ₅	0.5-50%
Metal oxides	1-15%

The resistor paste composition of the present invention is a mixture of the respective powders in the above proportions. Now, a process for producing the resistor paste of the present invention and a process for producing a thick film circuit using such a resistor paste, will be described.

An organic vehicle comprising an organic binder and a solvent, is added to the above resistor paste composition of the present invention, and the mixture is kneaded to obtain a paste. Such an organic binder includes ethyl cellulose, acrylic resins, ethylene-vinyl acetate copolymer resins and poly α -methylstyrene resins. Likewise,

the solvent includes α -terpineol, butylcarbitol acetate, butylcarbitol, 2,2,4-trimethylpentanediol-1,3-monoisobutylate, and diethylene glycol di-n-butyl ether. Further, a surfactant may be added as a dispersing agent.

Then, to form a conductor on a ceramic substrate such as a fired and solidified alumina substrate or on a glass ceramic substrate, a conductive paste such a Cu paste containing Cu as the main component, is formed in a prescribed circuit pattern by a method such as printing, followed by drying and then by firing at a temperature of from 800° to 1,000° C. for from 5 to 30 minutes in a non-oxidizing atmosphere such as a nitrogen atmosphere having an oxygen concentration of not higher than about 20 ppm. The preferred ranges of this firing conditions are from 880° to 920° C. for from 7 to 15 minutes. Then, the resistor paste of the present invention is printed at the predetermined portions for resistor, followed by drying and then by firing at a temperature of from 800° to 1,000° C. for from 5 to 30 minutes in the above-mentioned nitrogen atmosphere. The preferred ranges of this firing conditions are from 880° to 920° C. for from 7 to 15 minutes.

In the case of firing a multi-layer ceramic substrate all at once, ceramic green sheets for a ceramic substrate, having the above Cu paste and the resistor paste of the present invention already printed, are hot-pressed and laminated, followed by firing all at once at a temperature of from 800° to 1,000° C. for from a few minutes to a few hours in a non-oxidizing atmosphere such as the above nitrogen atmosphere, to obtain a multi-layer substrate.

To the resistor paste of the present invention, a coloring pigment such as a metal oxide or heat resistant inorganic pigment, may be incorporated for coloring in an amount of from 0 to 5%.

Further, from 0 to 5% of a nitrate, arsenic oxide, a sulfate, a fluoride or chloride may be added as a refining agent or a melting accelerator at the time of preparing the glass.

EXAMPLES

Starting materials for the glass powder of the present invention were mixed in the proportions as identified in Table 1, as calculated as oxides, and the proportions of the metal oxides are shown in Table 2. Each mixture was put into a platinum crucible heated under stirring at a temperature of from 1,350° to 1,500° C. for from 2 to 3 hours. Then, the product was pulverized in water or formed into flakes, and further pulverized by a pulverizer to an average particle size of from 0.5 to 6 μ m to obtain glass powder. Then, a powder of SnO₂ and/or SnO₂ having Sb doped in an amount of 5% as calculated as an oxide of Sb₂O₃, was adjusted to have an average particle size of from 0.01 to 5 μ m. Then, such glass powder and powder of conductive material were mixed in the proportions as identified in Table 1 to obtain a composition of the resistor paste of the present invention. Then, to such a resistor paste, an organic vehicle comprising ethyl cellulose as an organic binder and α -terpineol as a solvent, was added, and the mixture was kneaded to obtain a paste having a viscosity of 30 \times 10⁴ cps. Then, Cu was screen-printed as a conductor on a solidified alumina substrate in a prescribed circuit pattern, followed by drying and firing at 900° C for 10 minutes in a nitrogen atmosphere having an oxygen concentration of not higher than 20 ppm.

Then, at the prescribed portions for resistance, the above-mentioned resistor paste was screen-printed by a 200 mesh screen, followed by drying and firing at 900° C. for 10 minutes in a nitrogen atmosphere having an oxygen concentration of not more than 20 ppm. The film thickness after the firing was about 15 μm.

Thus, a circuit was formed on the ceramic substrate. With respect to this circuit, the resistivity, the temperature coefficient of resistivity (TCR) and the resistivity drift after storage at a high temperature, were measured. The results are shown in Table 1. It is evident from Table 1 that the resistor pastes of the present invention are excellent in the resistance characteristics and have adequate properties to be used as resistor pastes for forming thick film circuits.

As Comparative Examples, similar evaluations were conducted with respect to those other than the resistor pastes of the present invention. These are shown in Table 3.

The respective properties were measured by the following methods.

1) Resistivity and temperature coefficient of resistivity (TCR)

The resistivity values (R_{25} , R_{-55} and R_{125}) at 25° C., -55° C. and +125° C. were measured by an ohmmeter in a constant temperature tank, and the temperature coefficients of resistivity were calculated by the following formulas:

$$\text{Cold TCR} = \frac{R_{25} - R_{-55}}{R_{25} \times (25 - (-55))} \times 10^6 \text{ (ppm/}^\circ\text{C.)}$$

$$\text{Hot TCR} = \frac{R_{125} - R_{25}}{R_{25} \times (125 - 25)} \times 10^6 \text{ (ppm/}^\circ\text{C.)}$$

2) Resistivity drift after storage at a high temperature

The test sample was left to stand in a constant temperature tank of 150° C. for 100 hours, and the resistivity drift was calculated by the following formula:

$$\frac{R_{100h} - R_0}{R_0} \times 100 \text{ (%)}$$

In the above formula, R_{100h} = the resistivity after 100 hours, and R_0 = the initial resistivity.

TABLE 1

Sample No.	Examples								
	1	2	3	4	5	6	7	8	
Glass composition (%)	SiO ₂	20	30	45	30	20	25	15	14
	Al ₂ O ₃	10	0	0	10	5	0	18	0
	MgO	5	0	10	10	0	0	15	10
	CaO	20	10	10	15	10	2	5	0
	SrO	14	20	0	5	30	15	30	6
	Li ₂ O	0	0	2	0	0	0	0	0
	Na ₂ O	0	2	5	0	0	0	0	0
	K ₂ O	0	1	2	0	0	0	0	0
	Cs ₂ O	0	0	1	0	0	0	0	0
	PbO	0	3	0	0	0	0	0	0
	ZnO	5	0	0	15	0	0	0	0
	ZrO ₂	0	1	0	0	0	3	0	1
	TiO ₂	0	0	0	0	0	0	0	1
	B ₂ O ₃	25	21.5	18	9.5	30	35	11.5	8
	Ta ₂ O ₅	0	0	0	0	0.5	0	0	50
	Nb ₂ O ₅	0	10	0	0	0.5	0	0	0
	Constitution	Metal oxides	Present in Table 2						
Glass powder		40	50	55	40	25	48	70	45
Conductive									
SnO ₂		60	50	0	60	75	52	20	55
Average particle size (μm)	Sb-doped SnO ₂	0	0	45	0	0	0	10	0
	Glass powder	2.0	3.0	1.0	2.0	5.0	0.6	1.0	1.5
Properties	Powder of conductive material	1.0	0.1	1.0	0.05	0.5	5.0	0.1	0.2
	Resistivity (Ω/□)	16K	7M	1.5M	15K	7K	50K	12M	20M
	Hot TCR (ppm/°C.)	+70	+150	+30	-50	+100	+150	-120	+70
	Cold TCR (ppm/°C.)	+80	+120	+80	-70	+50	+160	-100	+80
	Resistivity drift (%)	+0.2	-0.8	+0.5	+0.2	+0.5	+0.2	-0.05	+0.8

Sample No.	Examples							
	9	10	11	12	13	14	15	
Glass composition (%)	SiO ₂	12	35	50	30	22	23	12
	Al ₂ O ₃	1	14	3	0	16	8	7
	MgO	0.5	1	25	2	32	0	0
	CaO	0.5	2	3	15	0	40	0
	SrO	58	14	0	30	0	0	58
	Li ₂ O	0.5	5	6	0	1	0	1
	Na ₂ O	0.5	1	0	0	0	2	2
	K ₂ O	0.5	3	0	0	0	0	5
	Cs ₂ O	0.5	0	0	0	0	0	1
	PbO	0.5	0	0	0	2	1	0
	ZnO	1	0	0	0	1	2	3
	ZrO ₂	0.5	0	0	0	2	1	1
	TiO ₂	0.5	0	0	0	1	2	1
	B ₂ O ₃	8	9	10	16	8	8	8
	Ta ₂ O ₅	0.8	0.5	1.5	2	0	1	0.5
	Nb ₂ O ₅	14.5	0.5	1	0	0	1	0

TABLE 1-continued

Constitution	Metal oxides	Present in Table 2						
	Glass powder	38	45	47	38	45	52	35
Conductive	SnO ₂	62	0	53	62	55	48	65
	Sb-doped	0	55	0	0	0	0	0
	SnO ₂							
Average particle size (μm)	Glass powder	2.2	6.0	3.0	2.6	3.2	3.0	2.7
	Powder of conductive material	0.5	1.2	0.8	0.2	1.0	0.5	1.0
Properties	Resistivity (Ω/□)	115K	150K	480K	3K	120K	620K	30K
	Hot TCR (ppm/°C.)	+100	+180	-130	-50	-80	-120	-250
	Cold TCR (ppm/°C.)	+120	+150	-160	-30	-100	-150	-230
	Resistivity drift (%)	-0.7	+1.0	+0.8	+0.2	-0.8	-0.7	+1.0
		Examples						
	Sample No.	16	17	18	19	20	21	22
Glass composition (%)	SiO ₂	15	15	20	17	12	15	17
	Al ₂ O ₃	12	10	10	0	20	18	13
	MgO	0	20	5	0	20	2	0
	CaO	0	2	1	10	0	30	0
	SrO	45	5	12	20	0	0	40
	Li ₂ O	0	0	10	0	8	0	0
	Na ₂ O	0	0	0	0	0	0	0
	K ₂ O	1	0	0	10	0	0	0
	Cs ₂ O	0	10	0	0	0	0	0
	PbO	0	5	10	8	0	0	2
	ZnO	2	0	0	10	2	0	0
	ZrO ₂	1	5	2	4	8	10	0
	TiO ₂	0	5	1	2	0	0	0
	B ₂ O ₃	9	10	10	11	10	10	8
	Ta ₂ O ₅	0	3	1	0	0	3	0
	Nb ₂ O ₅	0	0	1.5	0	1	2	0
Constitution	Metal oxides	Present in Table 2						
	Glass powder	32	50	40	45	41	40	37
Conductive	SnO ₂	68	50	60	55	59	60	63
	Sb-doped	0	0	0	0	0	0	0
	SnO ₂							
Average particle size (μm)	Glass powder	4.0	3.5	2.5	1.5	3.0	2.6	3.2
	Powder of conductive material	0.5	0.8	1.5	2.0	1.0	0.1	0.8
Properties	Resistivity (Ω/□)	18K	600K	15K	320K	20K	25K	85K
	Hot TCR (ppm/°C.)	-80	-20	+180	-100	+80	+90	-150
	Cold TCR (ppm/°C.)	-70	-40	+160	-80	+70	+100	-180
	Resistivity drift (%)	+0.6	-0.5	-0.8	+0.7	+1.0	+0.6	+0.8
		Examples						
	Sample No.	23	24	25	26	27	28	29
Glass composition (%)	SiO ₂	17	20	12	15	12	15	12.5
	Al ₂ O ₃	3	5	1.5	3	0	8	0.5
	MgO	0	6	10	8	12	5	0
	CaO	0	6	10	2	4	5	0
	SrO	50	6	10	10	0	8	16
	Li ₂ O	0	1	2	0	0	0	0
	Na ₂ O	0	1	3	2	0	0.5	0
	K ₂ O	0	1	1	1	0	0.5	0
	Cs ₂ O	0	1	0	2	0	0	0
	PbO	5	2	0	2	0	0	0
	ZnO	5	20	5	5	1	1	0
	ZrO ₂	0	2	0	1.5	0	0	0
	TiO ₂	10	3	0	2.5	0	0	0
	B ₂ O ₃	9.5	8	40	12	8	8	8
	Ta ₂ O ₅	0	1	3	15	55	0	18
	Nb ₂ O ₅	0	0	2	1	0	30	35
Constitution	Metal oxides	Present in Table 2						
	Glass powder	47	35	45	40	50	46	50
Conductive	SnO ₂	53	0	55	60	50	54	50
	Sb-doped	0	65	0	0	0	0	0
	SnO ₂							
Average particle size (μm)	Glass powder	3.5	3.0	2.5	2.7	4.0	1.0	0.5
	Powder of conductive material	0.01	0.1	1.0	2.5	1.4	1.5	5.0
Properties	Resistivity (Ω/□)	550K	2.5K	960K	200K	35M	15M	85M
	Hot TCR (ppm/°C.)	-200	-100	-250	-180	-220	-150	-250
	Cold TCR (ppm/°C.)	-220	-120	-200	-170	-200	-180	-220

TABLE 2-continued

Sample No.	Examples																					
	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44
WO ₃	0	0	0	2	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Bi ₂ O ₃	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
CeO ₂	0	0	0	0	0	0	2	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
CoO	0	0	0	2	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Cr ₂ O ₃	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Sb ₂ O ₃	0	0	0	0	0	0	0	0	0	0.5	5	15	11	0	0	0	0	0	0	0	0	0
In ₂ O ₃	0	0	0	0	0	0	0.5	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
SnO ₂	0	0	0	0.5	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0

Metal Oxides (wt %)	Sample No.	Examples																				
		23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43
NiO		0	0	0	0	0	0	0	0	0	0	0	0	0	0.1	5	10	6	4	5	0	0
Fe ₂ O ₃		0	0	0	0	0	0	0	0	0	0	0	0	0	0.5	1	0.5	0	0	0	0	0
CuO		0	0	0	0	0	0	0	0	0	0	0	0	0	0.1	1	0.5	0	0	0	0	0
MnO		0	0	0	0	0	0	0	0	0	0	0	0	0	0.1	1	3	0	0	0	0	0
MoO ₃		0	0	0	0	0	0	0	0	0	0	0	0	0	0.1	1	0.5	0	0	0	0	0
WO ₃		0.5		0	0	0	0	0	0	0	0	0	0	0	0.1	1	0.5	0	0	0	15	0
Bi ₂ O ₃		0	17	0	0	0	0	0	0	0	0	0	0	0	0.1	1	0.5	0	0	0	0	2
CeO ₂		0	0	0.5	18	0	0	0	0	0	0	0	0	0	0.1	1	0.5	0	0	0	0	0
CoO		0	0	0	0	8	19	0	0	0	0	0	0	0	0.1	1	0.5	0	0	0	0	0
Cr ₂ O ₃		0	0	0	0	0	0	10	20	0	0	0	0	0	0.1	1	0.5	0	0	0	0	0
Sb ₂ O ₃		0	0	0	0	0	0	0	0	0	0	0	0	0	0.1	3	2	0	0	0	0	0
In ₂ O ₃		0	0	0	0	0	0	0	0	0.5	5	20	0	0	0.1	1	0	0	0	0	0	0
SnO ₂		0	0	0	0	0	0	0	0	0	0	2	5	20	0.1	1	0	0	0	0	0	0

TABLE 3

	Sample No.	Examples		
		1	2	3
Glass composition (%)	SiO ₂	30	20	30
	Al ₂ O ₃	10	15	0
	MgO	0	5	5
	CaO	15	25	15
	SrO	20	10	15
	Li ₂ O	0	0	0
	Na ₂ O	0	2	2
	K ₂ O	0	5	0
	Cs ₂ O	0	0	0
	PbO	0	0	5
	ZnO	5	0	0
	ZrO ₂	0	2	0
	TiO ₂	0	1	0
	B ₂ O ₃	20	15	28
	Ta ₂ O ₅	0	0	0
	Nb ₂ O ₅	0	0	0
Constitution	Metal oxides	0	0	0
	Glass powder	10	40	50
	<u>Conductive material</u>			
	SnO ₂	90	60	50
Average particle size (μm)	Sb-doped SnO ₂	0	0	0
	Glass powder	3.0	2.0	1.0
	Powder of conductive material	1.0	1.0	0.5
Properties	Resistivity (Ω/□)	3K	30K	700K
	Hot TCR (ppm/°C.)	-2000	-1200	-800
	Cold TCR (ppm/°C.)	-2200	-1250	-900
	Resistivity drift (%)	+150	-2.0	+5.0

The resistor paste of the present invention can be fired in a non-oxidizing atmosphere such as a nitrogen atmosphere and is capable of forming a highly reliable stabilized resistor on a ceramic substrate, and it is particularly excellent in the resistivity drift properties after storage at a high temperature.

What is claimed is:

1. A composition of matter consisting essentially of an inorganic component which consists essentially of from 20 to 70 wt % of glass powder and from 30 to 80 wt % of a powder selected from the group consisting of SnO₂ powder, Sb-doped SnO₂ powder and a mixture thereof, wherein said glass powder consists essentially of from 12 to 50 wt % of SiO₂,

from 0 to 20 wt % of Al₂O₃, from 0 to 40 wt % of MgO, from 0 to 40 wt % of CaO, from 0 to 60 wt % of SrO, from 16 to 60 wt % of MgO+CaO+SrO, from 0 to 10 wt % of Li₂O+Na₂O+K₂O+Cs₂O, from 0 to 10 wt % of PbO, from 0 to 20 wt % of ZnO, from 0 to 10 wt % of ZrO₂+TiO₂, from 8 to 40 wt % of B₂O₃, either from 0.5 to 50 wt % of Ta₂O₅, or from 0.5 to 45 wt % of Nb₂O₅, or from 0.2 to 60 wt % of Ta₂O₅+Nb₂O₅, and from 0.1 to 20 wt % of Sb₂O₃.

2. A composition of matter consisting essentially of an inorganic component which consists essentially of from 20 to 70 wt % of glass powder and from 30 to 80 wt % of a powder selected from the group consisting of SnO₂ powder, Sb-doped SnO₂ powder and a mixture thereof, wherein said glass powder consists essentially of from 12 to 50 wt % of SiO₂, from 0 to 20 wt % of Al₂O₃, from 0 to 40 wt % of MgO, from 0 to 40 wt % of CaO, from 0 to 60 wt % of SrO, from 16 to 60 wt % of MgO+CaO+SrO, from 0 to 10 wt % of Li₂O+Na₂O+K₂O+Cs₂O, from 0 to 10 wt % of PbO, from 0 to 20 wt % of ZnO, from 0 to 10 wt % of ZrO₂+TiO₂, from 8 to 40 wt % of B₂O₃, either from 0.5 to 50 wt % of Ta₂O₅, or from 0.5 to 45 wt % of Nb₂O₅, or from 0.2 to 60 wt % of Ta₂O₅+Nb₂O₅, and from 0.1 to 20 wt % of MnO.

3. A composition of matter consisting essentially of an inorganic component which consists essentially of from 20 to 70 wt % of glass powder and from 30 to 80 wt % of a powder selected from the group consisting of SnO₂ powder, Sb-doped SnO₂ powder and a mixture thereof, wherein said glass powder consists essentially of from 12 to 50 wt % of SiO₂, from 0 to 20 wt % of Al₂O₃,

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from 0 to 40 wt % of MgO,
 from 0 to 40 wt % of CaO,
 from 0 to 60 wt % of SrO,
 from 16 to 60 wt % of MgO+CaO+SrO,
 from 0 to 10 wt % of Li₂O+Na₂O+K₂O+Cs₂O, 5
 from 0 to 10 wt % of PbO,
 from 0 to 20 wt % of ZnO,

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from 0 to 10 wt % of ZrO₂+TiO₂,
 from 8 to 40 wt % of B₂O₃,
 either from 0.5 to 50 wt % of Ta₂O₅, or
 from 0.5 to 45 wt % of Nb₂O₅, or
 from 0.2 to 60 wt % of Ta₂O₅+Nb₂O₅, and
 from 0.1 to 20 wt % of Sb₂O₃+MnO.
 * * * * *

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